

Applicants'

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ACCESSION NUMBER:

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TITLE:

Reactions with indole derivatives. XXVIII. Alkylation of camptothecin intermediates. New

approach to camptothecin

AUTHOR (S):

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CORPORATE SOURCE: Ger.

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Journal German

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For diagram(s), see printed CA Issue. The total synthesis of camptothecin was improved by using open-chain intermediates I, II, III, and IV to prep. 7-chlorodeoxycamptothecin (V), which improved the yield previously obtained during alkylation of VI. GI Several improvements of the previous synthesis were also described, e.g.,

the chlorination of the precursor of I.

IT

RL: SPN (Synthetic preparation); PREP (Preparation)

(prepn. of) 57182-95-3 CAPLUS

14H-Pyrano[3',4':6,7]indolizino[1,2-b]quinolin-14-one, RN 11-chloro-4-ethyl-1,12-dihydro- (9CI) (CA INDEX NAME) CN

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